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#### SEARCH REQUEST FORM

Scientific and Technical Information Center

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Requester's Full Name:	ZARA E	Examiner # : 77512 Da	nte: 12/30/02
Art Unit: 1630 Phone Nu	mber 30 6 - 5 8 20	Serial Number: () 9 /	915,814
Mail Box and Bldg/Room Location:			
If more than one search is submitt	************	*********	*****
Please provide a detailed statement of the se Include the elected species or structures, key utility of the invention. Define any terms the known. Please attach a copy of the cover sho	words, synonyms, acronymate may have a special mear eet, pertinent claims, and al	ns, and registry numbers, and comb ning. Give examples or relevant cit ostract.	oine with the concept or 🗀
Title of Invention:	5 to t	1 ) C Estima	ele-
Inventors (please provide full names):	Bester	etd	200
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Earliest Priority Filing Date: 7	126/01	<u> </u>	
*For Sequence Searches Only* Please include	all pertinent information (pa	rent, child, divisional, or issued paten	t numbers) along with the
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STAFF USE ONLY	Type of Search	Vendors and cost wher	e applicable
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Searcher Location:	Structure (#)	Questel/Orbit	
Date Searcher Picked Up: 12/03	Bibliographic	Dr.Link	
Date Completed: 13/03	Litigation	Lexis/Nexis	
Searcher Prep & Review Time:	Fulltext	Sequence Systems 106	
Clerical Prep Time:	Patent Family	WWW/Internet	<u> </u>
O-Use Times	Other	Other (specify)	

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January 3, 2003, 03:04:20; Search time 6400 Seconds (without alignments)
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TITLE JOURNAL	AUTHORS	REFERENCE			ORGANISM	SOURCE	KEYWORDS	VERSION	ACCESSION	DEFINITION	Locus	AX113509	RESULT 1
Patent: WO 0126664-A 1 19-APR-2001;	Mitchell, G.A. and wang, S.F.	1 (bases 1 to 3804)	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Howo.	Eukaryota; Metazoa; Chordata; Cranidia; Veriebraia, Horocome,	Homo sapiens	human.	•	AX113509.1 GI:13939/2/	AX113509	Sequence 1 from Patent WOUL20004.	IIINA IIICUI	2004	

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Scoring table: US-09-915-814-3 3804 1 cttcttgtaagagagtgcta.....taaataaaagtatttaatta 3804 IDENTITY\_NUC
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### ALIGNMENTS

AUTHORS TITLE JOURNAL	REFERENCE	ORGANISM	SOURCE	KEYWORDS	VERSION	ACCESSION	DEFINITION	LOCUS	RESULT 1 AX113509
Mitchell,G.A. and Wang,S.P. Hormone-sensitive lipase activity mediated male infertility Patent: WO 0126664-A 1 19-APR-2001;	<pre>bukaryota; Metazoa; Chordata; Crantata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 3804)</pre>	Homo sapiens	human.	•	AX113509.1 GI:13939727	AX113509	Sequence 1 from Patent WO0126664.	AX113509 3804 bp mRNA linear PAT 01-MAY-2001	

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1.141 CAGACACTACCAGGATACAGCCTCAAGGCTCATCCACACATGGACCTGCGCACAATGAC 1	4g 4g
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.021 CACGATGGGTGGAATGGTGGCCCAC	da An
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721 AGCTCAACAAGAAGCTGAATCAACACCTGCGGCCCAGGCTAAACCTGGAGCCAAAAAGGGA 780	. Qy
661 GCCAGCATTGAGACAAAGACATGTAGCCCAGCCAGGCCTGGGCCAGGAGAGACCACCTCC 720	p S
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220	2161 ACGGAGCCTGGAGCTGTGGCCGCGCCCCCAGCAGCACCCCGCTCGCGGTCCCTGATAGT 2	D Qy
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100	2041 CATCTCACCCCACTGGCCCACACAGGCCCTGGGCCCGTCCTCGTCAGGCTCATCTCCTA 2	Db Qy
040	1981 CAGCCTGCCACCCGAAGCCTTTGAGATGCCACTGACTGCCGACCCCACGCTCACGGTCAC 2	ОУ
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3481 AGACGGGGGCTGCGGGGGGGCGACACTAAAAGCCTGTTGTTCCCATCTGCGCCGGCC 	Qy 3,	
3421 CCTCGTCCTCACTCCTCCCGCCGGAGCCGGGCCGAGCGGGGAGACGGGGGCTGCGG 	Qy 3, Db 3,	
3361 CCTAGCGGCGCTGTGCCGCGAGACGCGCCAGGCCGCAGAGCTGTGCGTGGAGCGCA 	Qy 3.	
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Molecular cloning, genomic organization, and expression of a testicular isoform of hormone-sensitive lipase Genomics 35 (3), 441-447 (1996)
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/protein_id="AAC50666.1"
/db_xref="GI:1488677"
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/db_xref="taxon:9606"
/chromosome="19"
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Lund University, P.O. Box 94,
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                                                                                                                                                                                                                                                                           Direct Submission
Submitted (02-MAR-1993) Cecilia Holm,
University of Lund, PO. Box 94, Lund,
On Jul 13, 1995 this sequence version
Location/Qualifiers
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Homo sapiens male adult DNA.
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             /gene="LIPE"
/noce="600-120-154"
join(632. 1147,1148. .1238,1239. .1384,1385. .1570,
1571. .1865,1866. .2093,2094. .2270,2271. .2695,2696.
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                                                                                                   'number=1
                                                                                                                  /product="hormone-sensitive lipase"
/note="G00-120-154"
                                                                                                                                                     /gene-"LIPE"
                                                                                                                                                                   /cell_type="lymphocyte"
/dev_stage="adult"
|. .1147
                                                                                                                                                                                                                       sex="male"
                                                                                                                                                                                                                                                   /organism="Homo sapiens"
/db_xref="taxon:9606"
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PLTADPTLTTTTISPDLAHTGFGPVLVRLISYDLREGQDSEELSSLIKSNGQRSLELWP
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                   Score 2650.4;
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3245 GCGTGCGCGCTGGACCCCATGCTGGACGACTCCGGTCATGCTCGCGCGGCGACTGCGCAAC 3304	Qy Db	Qy 2165 AGCCTGGAGCTGTGGCCGCGCGCCCCAGCAGGCACCCCGCTCGCGGTCCCTGATAGTGCAC 2224	
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ICCASCASGETSCACACAGATSCCCCTCTACTCCTCACCCATAGTCAAAACCCCTTC		Qy 2045 TCACCCCCACTGGCCCACACAGGCCCTGGGCCCGTCCTCGTCAGGCTCATCTCCTATGAC 2104	
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2945 GACACCCCGAGATGTCGCTGTCAGCTGAGACACTTAGCCCCTCCACACCCTCCGATGTC 3004	Qy Db	Qy 1865 ATCACACAGAACCTGGACGTGCACTTCTGGAAAGCCTTCTGGAACATCACCGAGATGGAA 1924	
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GCCTCCTTGGCTCAACAGGGAACGAATCTGCCTTGCGGGGACAGTGCAGGCGGAAC 	Qy Db	Qy 1325 CCGGCCCTGGGCCGCCTGCGTGGGGTGTGGCGCACCTCTTTGACCTGGACCCAGAGACACCG 1384	
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TICCACGGCGGTGGCTTTGTGGCCCAGACCTCCAGAICCCACGAGCCCTACCTCAGAGCCTACTCAGAGCCTACCTCAGAGCCTACCTCAGAGCCCTACCTCAAGAGCTCCACGAGCCCCACGAGCCCTACCTCAAGAGCTCCACGAGCCCTACCTCAAGAGCCTACAGAGCCCTACCTCAAGAGCCCACGAGCCCTACCTCAAGAGCCCACGAGCCCTACCTCAAGAGCCCACGAGCCCTACCTCAAGAGCCACGAGCCCTACCTCAAGAGCCCACGAGCCCTACCTCAAGAGCCCACGAGCCCTACCTCAAGAGCCTACAGAGCCCTACAGAGCCCTAGAGAGCCCTACAGAGCCCTAGAGAGCCCTACCTCAAGAGCCAAGAGCCCTACAGAGCCCTACAGAGCCCTACCTCAAGAGCCCTACAGAGCCCTACCTCAAGAGCCCTACCTA	Db QY	Qy 1145 CACTACCAGGATACAGGCTCAAGGCTCACAACATGGACCTGCGCACAATGACACAG 1204	
AGCCTGGAGCTGTGGCCGCCCCCCCAGCAGGCACCCCGCTCGCGGTCCCTGATAGTGCAC	0; Db	Best Local Similarity 99.8%; Pred. No. 0; Matches 2654; Conservative 0; Mismatches 6; Indels 0; Gaps	

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                                                                              Direct Submission
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Biology, Lund University, P.O.
                                                                                                                            Stenson Holst, L., Langin, D., Laurell, H.,
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                                                                  Location/Qualifiers
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Eutheria;
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                                                                                 Stenson Holst, Cell and Molecular D. Box 94, Lund, S-22100, Sweden
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Query Match
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Matches 2740; Conserv
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                                                                                                                                  CACCTTACATCCAAAGGGTGCTGCTCACTCAACAGGAAGCTGCCTCCCAGCAGGGACCTG
                                                                                                                                                                                                                                                                                                    AACAAAATCTGCTTCACAAGAGGAATTTCTTGCCCCACAGAAGCCCGCACCACAGAAT
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                                                                               CATTCCCAAGCCACAAAGTGCACTCGGATCAACAGGAAGCCACCTCCCAGAATGGACCTG
                                                                                                                                                                                                                                     AACAATCTACTCTGCTCCAGAAACTTCTCACCCCATTAGCCTTCCCTGTACCACAGCAAT
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nilarity 73.0%;
Conservative
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/protein_id="AAC52771.1"
/db_xref="GI:1488679"
/da_xref="GI:1488679"
/da_xref
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 -GGGCAGAAGGATGAAACCTAGGAGACCAATTTCTTTCACAAGGGAAA
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Pred. No. 0;
0; Mismatches
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OM nucleic - nucleic search, using sw model
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Maximum Match 100%
Listing first 45 summaries
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Maximum DB seq length: 200000000
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Searched:
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Database
                                                          Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.
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17167.538 Million cell updates/sec
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Gapop 10.0 , Gapext 1.0
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1: /SIDS2/gcgdata/c
2: /SIDS2/gcgdata/c
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                                                                                                                                                                                                 SIDS2/gcgdata/geneseq/geneseqn-embl/NA1991.DAT:

SIDS2/gcgdata/geneseq/geneseqn-embl/NA1991.DAT:

SIDS2/gcgdata/geneseq/geneseqn-embl/NA1992.DAT:

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SIDS2/gcgdata/geneseq/geneseqn-embl/NA1994.DAT:

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SIDS2/gcgdata/geneseq/geneseqn-embl/NA1999.DAT:

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/SIDS2/gcgdata/geneseq/geneseqn-embl/NA1981.DAT:*
/SIDS2/gcgdata/geneseq/geneseqn-embl/NA1982.DAT:*
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/SIDS2/gcgdata/geneseq/geneseqn-embl/NA1986_DAT:*
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SUMMARIES
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Result		Query				•
NO.	Score	Match	Match Length DB	DB	ID	Description
_	3804	100.0	3804	22	AAD05012	Human mRNA encodin
2	3227.8	84.9	3231	24	ABK87218	Human lipase, horm
ω	1152.8	30.3		24	ABK87217	Human Lipase, horm
4	348.4	9.2		24	ABQ25066	Oligonucleotide to
ი 5	348.4	9.2		24	ABQ25067	Oligonucleotide to
6	307	8.1		22	AAI43422	Probe #12108 used
7	295	7.8		22	AAI56402	Probe #25088 used
c 8	285.6	7.5		24	ABQ25068	Oligonucleotide to
9	285.6	7.5		24	ABQ25069	Oligonucleotide fo
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#### ALIGNMENTS

RESULT 1 AAD05012

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AAD05012 standard; mRNA; 3804 BP
                                                                                                                                                                                               Human; hormone sensitive lipase; contraceptive; antisense-therapy; inhibitor; testicular isoform; male fertility; ss.
                                                                                                                                                                                                                            Human mRNA encoding testicular isoform of hormone sensitive lipase
                                                                                                                                                                                                                                                                      AAD05012;
Mitchell GA,
                                                        13-OCT-2000; 2000WO-CA01228.
                                                                                               WO200126664-A2
                                                                                                                                                                                                                                                  17-JUL-2001 (first entry)
                                                                             19-APR-2001.
                                                                                                                                                                             Homo sapiens.
                 (HOPI-) HOPITAL SAINTE-JUSTINE.
                                      14-0CT-1999;
 Wang SP;
                                      99CA-2286451.
                                                                                                                  /*tag= a
/product= "Human hormone sensitive lipase"
/EC_number= "3 1.1.3"
                                                                                                                                                Location/Qualifiers
278..3508
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Best Local Sim.
Matches 3144;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The present sequence is a mRNA encoding the testicular isoform of human hormone-sensitive lipase (E.C. 3.1.1.3). The present invention relates to a method of inhibiting male fertility which involves inhibition of the activity of the hormone-sensitive lipase. The hormone sensitive lipase, especially the testicular isoform from humans is useful for screening compounds that modulate or inhibit male fertility. Inhibitors of hormone sensitive lipase, especially a fully defined antisense molecule are useful for inhibiting fertility in a male animal. The method provides means for inhibiting fertility in genetically modified organisms to prevent the dissemination of genetic modifications into wild-type populations. The method is also used to identify a condition of male infertility caused by hormone sensitive lipase deficiency.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Sequence
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                 ACAGGAAGCTGCCTCCCAGCAGGGACCTGGGCTAGGAAAAGAATCTATAACTCAACAGGA
                                                      TGCTGAATCCCAGAAGGAACCTAGAGCCCCAACAAAAATCTGCTTCACAAGAGGAATTTCT
                                                                                                                                                                                                                          GCCAGAAAAGACACCCATAGCCCAGCCAGAATCGAAGACTCTGCAGGGATCCAATACCCA
ACAGGAAGCUGCCUCCCAGCAGGGACCUGGGCUAGGAAAAGAAUCUAUAACUCAACAGGA
                                                                                                               UGCUGAAUCCCAGAAGGAACCUAGAGCCCCAACAAAAAUCUGCUUCACAAGAGGAAUUUUCU
                                                                                                                                                                      ACAGAAGCCUGCUUCAAACCAAAGACCCCUCACCCAGCAGGAGACCCCCUGCACAACAUGA
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)B; AAE01154.
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                                                            GCTCACCGCCGACTTCCTCCGGGAGTATGTCACGCTGCATAAGGGATGCTTCTATGGCCG
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                                                                                     ACGCTCCAGCCAGGGTGCCACACAGATGCCCCTCTACTCCTCACCCATAGTCAAGAACCC
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       GTTGTAAATAAAAGTATTTAATTA
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                                                         GCACACACCGGUCACCGAGACGGCUGGACCUGCACGCCACCGCUGCCUUUUUGCUGCUG
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male sterility; single nucleotide polymorphism; SNP; chromosome 19q13.1-q13.2; gene; ss.
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replace (280,A)
/*tag= b
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/standard_name= "Single nucleotide polymorphism /note= "Polymorphic site (PS)" replace (2453,A) /*tag= p
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/note= "Polymorphic site (PS)"
replace (1677,A)
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/*tag= i
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/*tag= h
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/*tag= g
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278 ATGGAGCCAGGTTCTAAGTCAGTGTCTAGGTCAGACTGGCAACCTGAACCACCACCAGAGG 337

Matches Query Match Best Local

Similarity

84.9%; 99.9%;

0;

Pred. No. 0; ); Mismatches Score

2;

Indels

0;

Gaps

0

Length

3227. . 8

Sequence 3231

BP;

693 A; 1080 C; 906 G; 552

<del>,</del> DΒ

0 other;

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nucleotide sequence which comprises lipase, hormone sensitive (LIPE) isogenes. The invention is useful in screening for drugs targeting LIPE isogenes that are useful for treating obesity and male sterility. The methods of the invention are useful for improving the efficiency and reliability of several steps in the discovery and development of drugs for treating diseases associated with LIPE activity. The polynucleotide is useful in studying the expression and function of LIPE, and in expressing LIPE protein for use in screening for candidate drugs to treat diseases related to LIPE activity. It is also useful in studying the effect of the variation on the biological activity of LIPE as well as on the binding affinity of candidate drugs targeting LIPE for the treatment of obesity and male sterility. The invention is useful for studying the expression of LIPE 1sogenes in vivo, for in vivo screening and testing of the treatment are treatment as a series of the control of
                                                                 drugs targeted against LIPE protein, and for testing the efficacy of therapeutic agents and compounds for treating obesity and male sterility in a biological system. The present nucleic acid sequence represents the coding sequence of the human LIPE gene located on chromosome 19q13.1-q13.2. This sequence encodes the human LIPE protein of the
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Novel genetic variants of Lipase, Hormone-Sensitive isogenes, for improving efficiency and reliability in drug development i treating diseases associated with LIPE activity, e.g. obesity
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/note= "Polymorphic site (PS)"
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/note= "Polymorphic site (PS)"
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/note= "Polymorphic site (PS)"
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/*tag=
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/note= "Polymorphic site 9 (PS9)"
replace (20676,T)
/*tag= n
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4084..29321
/*tag= b
               replace (22783,
/*tag= p
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/note= "Polymorphic site
replace (5041,T)
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4967..20290
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                                                                        20827..22810
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/note= "Polymorphic
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replace (4612,A)
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/note= "Polymorphic site
replace (4519,T)
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replace (4381,C)
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replace (4717,C)
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<pre>/*eplace (25714,T) /*tag= ak /*standard_name= "Single nucleotide polymorphism (SNP)"</pre>	/ replace (25659,A) /note= "Polymorphic site 19 (PS19)" replace (25659,A) /*tag= aj /*standard_name= "Single nucleotide polymorphism (SNP)" /note= "Bolymorphic site 20 (BS20)"	18 (PS18)".	ngle	/number= 7 /number= 7 2557225748 /*tag= ag	/note= "Polymorphic site 17 (PS17)" 2474524972 /*tag= ae /number= 7 249732571	/"Lag= ac /number= 6 replace (24064,G) /*tag= ad /standard_name= "Single nucleotide polymorphism (SNP)"	<pre>/*tag= ab /standard_name= "Single nucleotide polymorphism (SNP)" /note= "Polymorphic site 16 (PS16)" 2396024744</pre>	2366523959 /*tag= aa /number= 6 replace (23727,A)	replace (23608,A)  /*tag= z  /*tag= z  /*tandard_name= "Single nucleotide polymorphism (SNP)"  /note= "Polymorphic site 15 (PSI5)"	/Standard_hame= "Single nucleotide polymorphism (SNP)" /note= "Polymorphic site 14 (PS14)" 2352823664 /*tag= y /numbor= S	13 (PS13)"	P)	/number= 4 23341.23527 /*tag= v	: _ 2	T)	. H	/number= 3 22903.23012 /*tag= r	
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GGCGTGCGCGTTGGATTTTATGTTGGACGATTCGGTTATGTTCGCGCGGGGGATTGCGTAA 178

CCTGGGCCAGCCGGTGACGCTGCGCGCGTGGTGGACCCTGCCGCACGGCTTCCTGACCCT 336

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 ${ t AGCGGCGTTGTCGCGAGACGCGTTAGGTCGTAGAGTTGTGCGTGGAGCGTATTCGTTT}$ 

Query Match Best Local Matches

Similarity

9.2%; 79.5%;

450;

Conservative

0;

Score 348.4; Pred. No. 8.9e 0; Mismatches

8.9e-66;

111;

Indels Length

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Gaps

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DB 24;

Sequence

BP;

97 A;

74 C; 238

G;

300 T; 0 other;

disclosure of the invention.

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This invention describes a novel method for determining the degree of CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a genomic sample of DNA. The sample is treated chemically to convert CC genomic sample of DNA. The sample is treated chemically to convert CC DNA that contains the target C is amplified to form a labeled amplicon. The amplicon is hybridised to two classes, each with at least one member, of oligonucletides and/or peptide-nucleic acid (PNA) oligomers CC and the degree of hybridisation to both classes is determined from the CC label on the amplicon. From the ratio of labels hybridised to the two CC classes of oligomers, the degree of methylation is calculated. The method is used: (i) for diagnosis and/or prognosis of side effects of CC is used: (i) for diagnosis and/or prognosis of side effects of CC therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders of the central nervous, cardiovascular, gastrointestinal and respiratory systems etc., particularly by detecting mutations or single nucleotide polymorphisms (SNP's); and (ii) for differentiation. The method allows the CC methylation status of many C residues to be determined simultaneously.

CC methylation status of many C residues to be determined simultaneously.

CC methylation status of many C residues to be determined simultaneously.

CC methylation status of many C residues to be determined simultaneously.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                           Determining the degree of cytosine methylation in genomic DNA, for diagnosis and prognosis, comprises selective hybridization amplicons from chemically treated DNA .
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05-SEP-2000;
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gastrointestinal; respiratory system;
SNP; cell differentiation; ds.
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ABN52192
                                                                                                                                                              ABN32626
                                                                                                                                                                                                                                                               SUMMARIES
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ; Search time 499 Seconds (without alignments) 17167.538 Million cell updates/sec
Human spliced tran
Human SNP oligonuc
Human SNP oligonuc
Rat spliced transc
Human SNP oligonuc
Human SNP oligonuc
Human SNP oligonuc
Mouse spliced transc
Mouse spliced transc
Mouse spliced transc
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O	C	C	C					C		O	O	C		O			C			O	C					a	O	ი	O	ი	a			C	
45	44	43	42	41	40	39	38	37	36	35	34	ω 3	32	31	30	29	28	27	26	25	24	23	22	21	20	19	18	17	16	15	14	13	12	11	10
25	•	•	25.2	•	•	•	•	•	•	•	•	•	•	٠			•	N	26	26	6	σ,	6	26.4	σ,	σ.	σ,	σ,	9	σ,	27	27	27.6	7	29
	•	•	0.7	•	•	•	•	•	•	•	•	•	•	•	•	•	•	•	•	0.7	•	•	•	•	•	•	•	•	•			•		•	
66	100	89	65	61	61	61	61	100	96	100	99	99	70	66	65	64	99	94	92	90	87	86	100	100	100	65	65	65	99	99	72	60	60	89	88
17	22	22	19	22	22	21	18	17	21	24	21	14	21	12	24	20	21	19	21	24	16	24	24	22	22	22	22	21	18	17	20	24	24	19	23
AAT97798	AAH41935	AAS32956	AAV41235	AAF23370	AAF74334	AAZ88282	AAT84492	AAT15244	AAZ57144	ABL74850	AAC98715	AAQ36997	AAA38176	AAQ12354	ABN57988	AAX88104	AAA36736	AAV38341	AAC11989	ABK36582	AAT22575	AAS98931	ABS20120	AAK45837	AAK19814	AAF23371	AAF74335	AAZ88283	AAT96968	AAT27118	AAX88105	ABN32825	ABN45730	AAV68382	AAS48737
Nucleic acid inhib	LASP-3	DNA encoding CARDI	embry	gene ]		SlpIII nucleotide	SlpIII gene sectio	Partial coding seq	Exemplary quadrupl	Corn tassel-derive	Human colon cancer	0	Primer used in ste	Europium-labelled	Mouse spliced tran	Synthetic antifree	MuV and MV fusion	Manic-depressive i	Human secreted pro	HCV DNA encoding H	Human gene signatu		Human genome-deriv			I gene	Repetitive protein	cle	Œ	м	reeze pro	spliced	spl	lone #8 fra	Pseudomonas aerugi

## ALIGNMENTS

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RESULT 1
ABN32626
ABN32626;
                                                                                                      Human spliced transcript detection oligonucleotide SEQ ID NO:5374.
                                                                                                               15-JUL-2002 (first entry)
                                                                                                                                 ABN32626 standard; DNA; 60
                                                                                                                                 ВP
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28-JUL-2000; 2000US-221607P 02-MAY-2001; 2001US-287724P Human; mouse; rat; splice transcript; detection; RNA transcript; splice variant; transcriptome; oligonucleotide library; ss. 20-JUL-2001; 2001WO-IB01903 07-FEB-2002. WO200210449-A2 Homo sapiens.

Shoshan A, (COMP-) COMPUGEN INC Wasserman A,

Mintz Ĺ Mintz Ļ Faigler

WPI; 2002-257383/30

New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription unit of

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AAL33801
ID AAL
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               CC transcription units that populate a genome. The library comprises CC several oligonucleotides, each capable of hybridising selectively to a CC the genome, which encodes one or more messenger RNA splice variants. CC the genome, which encodes one or more messenger RNA splice variants. CC biological sample, in expression profiling studies, in qualitatively or quantitatively characterising the corresponding transcriptome, and in CC quantitatively characterising the corresponding transcriptome, and in CC transcriptomes. The libraries may also be used as specialised mini CC transcriptomes. The libraries may also be used as specialised mini CC particular biological or pathological state, and so allowing the CC only expressed in specific tissue under a specific pathological condition; to detect developmental specific genes such as those genes condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering CC oligonucleotide sequences from rats, humans and mice, which are used in CC the exemplification of the present invention.

CC Specification, but was obtained in electronic format directly from WIPO CC at ftp. wipo.int/pub/published not seminances
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Query Match
Best Local Similarity
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Matches
                                                                                                                                                                                                                                                                         Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic; neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer; amyloid proteain; angiopoietin; apoptosis related protein; cadherin; cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor; complement related protein; cytochrome; kinase; cytokine; interferon; interleukin; G-protein coupled receptor; thioesterase; inflammation; multifactorial disease; autoimmune disease; infection; nervous system disease; ss.
(CURA-) CURAGEN CORP
                                            28-DEC-1999;
27-DEC-2000;
                                                                                                        28-DEC-2000; 2000WO-US35498
                                                                                                                                                                                                     WO200147944-A2
                                                                                                                                                                                                                                             Homo sapiens.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Human SNP oligonucleotide #7009.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             24-JAN-2002 (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        AAL33801 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          2621 AAGTGTGTCAGCGCCTATGCTGGTGCAAAGACGGAGGACCACTCCCAACTGAGACCAGAAA 2680
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Sequence 60
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              developmental-specific
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (sub-)transcriptome
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ftp.wipo.int/pub/published_pct_sequences
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     genome, useful for detecting tissue-, pathology-, and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AAGTGTGTCAGCGCCTATGCTGGTGCAAAGACGGAGGACCACTCCAACTCAGACCAGAAA 60
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      60;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              nt invention describes oligonucleotide libraries for detecting RNAs that populate a (sub-)transcriptome, where the nscriptome comprises messenger RNAs transcribed from multiple
                                     99US-0173419
2000US-0173419
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          BP;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     DNA;
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Pred. No.
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0.0012;
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Дb Ş

WPI; 2001-465210/50

Leach

28-DEC-1999; 27-DEC-2000;

2000US-0173419.

99US-0173419

(CURA-) CURAGEN CORP

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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             CC The present invention relates to oligonucleotides encoding polymorphic CC variants of proteins related to amylases, amyloid proteins, anglopoietin, CC apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes, CC histones, kinases, colony stimulating factors, complement related CC G-proteins, cytochromes, kinesins, cytokines, interferons, interleukins, CC one such oligonucleotide. The oligonucleotides and the present sequence is CC one such oligonucleotide. The oligonucleotides and treatment of CC by them may be used in the prevention, diagnosis and treatment of CC diseases associated with inappropriate expression of the proteins listed CC above. Disorders that may be prevented, diagnosed and/or treated include CC multifactorial diseases with a genetic component, such as autoimmune CC systemic lupus erythromatosus and Grave's diseases), diabetes, CC (e.g. cancers of the bladder, brain, breast, colon and kidney, CC corqanisms.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Matches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Query Match
Best Local
                                                                                                                                                                                                                                                                            complement related protein; cytochrome; kinesin; cytokinesin; intering interleukin; G-protein coupled receptor; thioesterase; inflammation; multifactorial disease; autoimmune disease; infection:
                                                                                                                                                                                                                                                                                                               Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic; neuroprotective; antimicrobial; gene therapy; vaccine; amylase; can amyloid protein; andiopoletin; apoptosis related protein; cadherin; cyclin; polymerase; oncogene; histone; kinase; colony stimulating f
                                                                                                                                                                   05-JUL-2001
                                                                                                                                                                                                 WO200147944-A2
                                                                                                                                                                                                                                Homo sapiens.
                                                                                                                                                                                                                                                               nervous system
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            1516 GGCCTACCTGGCTGCCCTCACCCAGCTCCGCGCTCTGGTCTACTACGCCCA 1566
                                                                                                                                                                                                                                                                                                                                                                                                 Human SNP oligonucleotide #7012.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                  AAL33804;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           AAL33804 standard; DNA; 51 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Sequence 51
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Claim 1; Page 3391; 4143pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      cancer, autoimmune
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Polymorphic nucleic acids encoding e.g. amylases, oncogenes and histones, useful for diagnosing and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Shimkets RA,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 1 GCCTACCTGCCTCACCCAGCTCCGCGCTCTGGTCTACTACGCCCA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Similarity
                                                                                                                                    2000WO-US35498
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                (first entry)
                                                                                                                                                                                                                                                            disease;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Α;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1.3%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  diseases and infections
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              24 C;
                                                                                                                                                                                                                                                            autoimmune disease; ss.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         11 G; 10 T; 0 other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Score 51; Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             DB 22;
0.099;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Length 51;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 51
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            0;
                                                                                                                                                                                                                                                                                                                                              cancer;
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RESULT 4
ABN28875
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Query Match
                                                                                                                                                                                                                                                                                                                                                                                           Human;
splice
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       by them may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate expression of the proteins listed above. Disorders that may be prevented, diagnosed and/or treated include multifactorial diseases with a genetic component, such as autoimmune diseases (e.g. rheumatoid arthritis, multiple sclerosis, diabetes, systemic lipus erythromatosus and Grave's disease), inflammation, cancer (e.g. cancers of the bladder, brain, breast, colon and kidney,
                                                                                                                                 WPI;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             proteins, cytochromes, kinesins, cytokines, interferons, interleukins, G-protein coupled receptors and thioesterases. The present sequence is one such oligonucleotide. The oligonucleotides and the peptides encoded
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           variants of proteins related to amylases, amyloid proteins, angiopoiet apoptosis related proteins, cadherin, cyclin, polymerase, oncogenes, histones, kinases, colony stimulating factors, complement related proteins, cytochromes, kinesins, cytokines, interferons, interleukins,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Polymorphic nucleic acids encoding {\rm e.g.} amylases, oncogenes and histones, useful for diagnosing and cancer, autoimmune diseases and infections -
                                                                     New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription a genome, useful for detecting tissue-, pathology-, and
                                                                                                                                                                                                                         28-JUL-2000;
02-MAY-2001;
                                                                                                                                                                                                                                                                                                                                                                                                                                         Rat spliced transcript detection oligonucleotide SEQ ID NO:1623
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   ABN28875 standard; DNA; 65 BP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       2160 AACGGAGCCTGGAGCTGTGGCCGCGCCCCCAGCAGGCACCCCGGCTCGCGGT
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Sequence 51 BP;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       The present
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Claim 1; Page 3392; 4143pp; English.
                                                          developmental-specific
                                                                                                                                                                                                                                                                       20-JUL-2001; 2001WO-IB01903
                                                                                                                                                                                                                                                                                                     07-FEB-2002
                                                                                                                                                                                                                                                                                                                                 WO200210449-A2
                                                                                                                                                                                                                                                                                                                                                             Rattus norvegicus
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     ABN28875
                                                                                                                                                                                             (COMP-) COMPUGEN INC.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                       15-JUL-2002
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         leukaemia),
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           _
                                                                                                                                  2002-257383/30
                                                                                                                                                                                                                                                                                                                                                                                           mouse; rat;
variant; tra
                                                                                                                                                               P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Similarity
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            diseases
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         invention relates to oligonucleotides encoding polymorphic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                     (first entry)
                                                                                                                                                                                                                         2000US-221607P
2001US-287724P
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              7 A;
                                                                                                                                                                                                                                                                                                                                                                                          at; splice transcript; detection;
transcriptome; oligonucleotide l:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1.3%;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          of the nervous system and an infection of pathogenic
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              21 C; 18 G; 5 T; 0 other;
                                                                                                                                                               Ą
                                                          genes
                                                                                                                                                               Mintz
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Score 51;
Pred. No.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Mismatches
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                                                                                                                                                               Mintz L,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0.099;
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                                                                                                                                                               Faigler
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    0;
                                                                                                                                                                                                                                                                                                                                                                                             n; RNA transcript;
library; ss.
                                                                        a transcription unit gy-, and
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Length 51
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   cyclins, polymerases, treating, e.g.
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Gaps

0;

The present invention describes oligonucleotide libraries for detecting

Example

1;

SEQ

IJ

1623; 47pp;

English

of

Shimkets RA,

Leach M;

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RESULT 5
AAL33802
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         δÃ
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Matches
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Best Local
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               messenger RNAs that populate a (sub-)transcriptome, where the (sub-)transcriptome comprises messenger RNAs transcribed from multiple transcription units that populate a genome. The library comprises several oligonucleotides, each capable of hybridising selectively to a set of messenger RNAs transcribed from a given transcription unit of the genome, which encodes one or more messenger RNA splice variants. The oligonucleotide libraries are useful for detecting mRNAs from a biological sample, in expression profiling studies, in qualitatively or quantitatively characterising the corresponding transcriptome, and in detecting RNA transcripts and splice variants of human or animal transcriptomes. The libraries may also be used as specialed minimal transcriptomes. The libraries may also be used as specialed minimal transcriptomes.
                                                                                                                                                                                                                                                                                               Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic; neuroprotective; antimicrobial; gene therapy; vaccine; amylase; cancer amyloid protein; angiopoietin; apoptosis related protein; cadherin; cyclin; polymerase; oncogene; histone; kinase; colony stimulating fact complement related protein; cytochrome; kinesin; cytokine; interferon; interreukin; G-protein coupled receptor; thioesterase; inflammation;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          3192
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          condition; to detect developmental specific genes; and to detect RNA transcripts and splice variants of a transcriptome of a patient suffering from a particular disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from rats, humans and mice, which are used in
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             the exemplification of the present invention. N.B. The sequence data for this patent did not specification, but was obtained in electronic at ftp.wipo.int/pub/published_pct_sequences.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                libraries to detect transcripts of a sub-transcriptome under a particular biological or pathological state, and so allowing the detection of tissue- and pathology-specific genes such as those genes only expressed in specific tissue under a specific pathological
                                                                                                                                                                                                                                                                                                                                                                                                                    Human SNP oligonucleotide #7010.
                                                                                                                                                                                                                                                                                                                                                                                                                                                       24-JAN-2002
                                                                                                                                                                                                                                                               nervous system disease;
                                                                                                                                                                                                                                                                                    multifactorial
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          AAL33802 standard;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             Sequence
                                                                              28-DEC-1999;
27-DEC-2000;
                                                                                                                               28-DEC-2000; 2000WO-US35498
                                                                                                                                                                 05-JUL-2001.
                                                                                                                                                                                                                                Homo sapiens.
                                               (CURA-) CURAGEN CORP
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            61
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            CGCTG
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              | Similarity
| 56; | Conser
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               65
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              Conservative
                                                                              99US-0173419
2000US-0173419
                                                                                                                                                                                                                                                                                                                                                                                                                                                       (first entry)
                                                                                                                                                                                                                                                                                   disease; autoimmune disease;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               13
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Pred. No. 0.13
0; Mismatches
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                                                                                                                                                                                                                                                                                    infection;
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format directly from
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RESULT 6
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Query Match
Best Local Similarity
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Polymorphic nucleic acids encoding e.g. amylases, cyclins, polymerases,
                                   WPI; 2001-465210/50
                                                                        Shimkets
                                                                                                                                            28-DEC-1999;
27-DEC-2000;
                                                                                                                                                                                                  28-DEC-2000; 2000WO-US35498
                                                                                                                                                                                                                                                                       WO200147944-A2
                                                                                                                                                                                                                                                                                                                                          interleukin; G-protein coupled receptor; thioesterase; inflammation;
multifactorial disease; autoimmune disease; infection;
nervous system disease; ss.
                                                                                                                                                                                                                                                                                                                                                                                           amyloid protein; angiopoietin; apoptosis related protein; cadherin; cyclin; polymerase; oncogene; histone; kinase; colony stimulating factor; complement related protein; cytochrome; kinesin; cytokine; interferon;
                                                                                                                                                                                                                                                                                                                                                                                                                            Immunosuppressive; immunostimulatory; antiinflammatory; cytostatic; neuroprotective; antimicrobial; gene therapy; vaccine; amylase; can amyloid protein; angiopoletin; apoptosis related protein; cadherin;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Human SNP oligonucleotide #7011.
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                                                                                                                                                                                                                                                                                                          Homo sapiens
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                                                                                                        (CURA-) CURAGEN CORP
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2000US-0173419
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                                                                      Leach M;
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Pred. No.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                    cancer;
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RESULT 7
ABN53647
Вb
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Query Match
Best Local
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02-MAY-2001;
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                                                                                                                                                     WPI; 2002-257383/30
                                                                                                                                                                                Shoshan A,
                                                                                                                                                                                                                                                                                                             07-FEB-2002
                                                                                                                                                                                                                                                                                                                                                                                                              Human;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               Sequence 50
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                                                                                                                                                                                                             (COMP-)
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    Local
                                                                                                                                                                                                                                                                                                                                                                                               mouse; rat; splice transcript; detection; RNA transcript;
variant; transcriptome; oligonucleotide library; ss.
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                                                                                                                                                                                                             COMPUGEN INC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               autoimmune diseases and infections
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2001US-287724P.
                                                                                                                                                                                                                                                                                  2001WO-IB01903
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   1.0%;
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                                                                                                                                                                                A,
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                                                                                                                                                                                Mintz E,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               ВP
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                                                                                                                                                                                Mintz L,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         0;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   Length 50;
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The present invention describes oligonucleotide libraries messenger RNAs that populate a (sub-)transcriptome, where

(sub-)transcriptome

comprises

messenger RNAs

transcribed

for

detecting multiple

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Example 1; SEQ

ID 26395; 47pp; English

New oligonucleotide libraries comprising oligonucleotides which selectively hybridize to mRNAs transcribed from a transcription a genome, useful for detecting tissue-, pathology-, and developmental-specific genes

unit of

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Title:
Perfect score:
Sequence:
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Maximum
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       DB
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Result No.

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SUMMARIES

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47.1 31.9 24.2 21.3 20.7 20.6

1809 2760 1084 826 872 905

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BC029961 Homo sapi AK019846 Mus muscu BM563641 AGENCOURT BI826568 603077016 BI827841 603074047 BI827706 603074127

REMARK COMMENT	REFERENCE AUTHORS TITLE JOURNAL	RESULT 1 BC029961 LOCUS DEFINITION ACCESSION VERSION KEYWORDS SOURCE ORGANISM		1ω4π 1	2 1 0	98	765	<b>4</b> ω κ	) <del>L.</del> C	987	400	ω N P (	c 17 c 18 c 19 6	4100	ω N P C	987
JRL: http://mgc.nci.nil o desk nail.nih.gov nt: Life Technologies, paration: Life Technolo syed by: The I.M.A.G.E y: Baylor College of Mc G. HGSC www.hgsc.bcm.tmc.edu/ .tmc.edu	Manmalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  E 1 (bases 1 to 1809)  S Strausberg, R.  Direct Submission L Submitted (06-MAY-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,	BC029961  1809 bp mRNA linear Homo sapiens, clone IMAGE:5171623, mRNA. BC029961 BC029961.1 GI:20455827 HTC. Homo sapiens. Homo sapiens. Homo sapiens.	ALIGNMENTS	73.2 12.4 600 12 473 12.4 588 12 469 12.3 469 10	12.6 952 12 BF537918 BF5379 12.5 882 12 BG868403 BG8684 12.5 913 13 B1654202 B16542	496 13.0 539 10 AW027306 AW0273 92.2 12.9 643 12 BF222934 BF2229	13.1 557 14 BM714935 BM7149 13.1 553 14 BM675647 BM65766 13.1 572 9 AA639729 AA6397	10.4 13.4 892 14 BQ953768 BQ9537 03.2 13.2 669 9 AA861626 AA86162 00.6 13.2 1036 13 BM560162 BM5601	21.4 13.7 741 12 BG872368 BG8723 20.4 13.7 524 10 AW195614 AW1956	48.2 14.4 555 9 AI798896 AI79889 38.6 14.2 635 9 AA601541 AA60154 534 14.0 534 12 BE855571 BE8555	558 14.7 985 12 BG023944 BG0239 56.2 14.6 653 9 AI393944 AI39394 55.8 14.6 601 9 AI582719 AI58271	16.0 691 13 BI818150 BI818181 14.9 569 10 AW269990 AW2699 14.9 588 14 BQ807836 BQ8078	625 13 BM310576 BM3105 668 14 BQ028786 BQ028786 736 9 A1953697 A195369 647 13 B1855956 B18559	17.6 1367 11 BC029301 BC0293 17.1 663 13 B1826036 B18260 17.0 645 13 B1825158 B18251	73.2 13.0 700 13 B1520440 B152016 73.6 19.1 773 13 B1520160 B15201 1.2 18.7 781 13 B1832284 B18322 697 18.3 767 9 A1337272 A133727	82.6 20.6 870 13 B1827559 B18275 64.8 20.1 871 13 B161175 B15611 757 19 9 787 13 B1830913 B18309

Gunaratne, P.H., Garcia, A.M., Lu, Yoon, V.S., Kowis, C.R., Lawrence, Richards, S., Gibbs, R.A.

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This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 488545(This clone has the following problem: no polyA-tail.
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/db_xref="taxon:9606"
/clone="IMAGE:5171623"
/tissue_type="Brain, adult me/clone_lib="NIH_MGC_119"
/lab_host="DH10B"
/note="Vector: pCMV-SPORT6"
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High-efficiency full-length cDNA cloning
Meth. Enzymol. 303, 19-44 (1999)
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Mus musculus (strain:C57BL/6J) adult male testis cDNA to mRNA, clone_lib:RIKEN_full-length enriched mouse cDNA library
5 (bases 1 to 2760)
Adachi, J., Aizawa, K.,
                                                                                                                                                                                              Functional annotation of a full-length mouse cDNA collection Nature 409 (6821), 685-690 (2001)
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AIRPFLOTLSIGLYSFGEHYKRNETGLSVTASSLFTGGRFAIDPELRGAEFERIIQNL
DVHFWKAFWNITEIEVLSSLANNASTTVRVSRLISLIPEPEAFEMFUTSDPRLTVTISPP
LAHTGPAPVLARLISYDLREGODSKVLNSLAKSEGPRLELRPRPHQAPRSRALVVHIH
GGGFVAOTSKSHEPYLKNWAOBLGVPJIFSIDYSLAPEAPFPRALEECFFAYCWAVKHC
DLLGSTGERICLAGDSAGGNLCITVSLRAAAYGVRVPDGIMAAYPVTTLOSSASPSRL
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HFDLDTETPANGYRSLVHTARCCLAHLLHKSRYVASNRKSIFFRASHNLAELEAYLAA
LTQLRAMAYYAQRLLTINRPGVLFFEGDEGLTADFLQEYVTLHKGCFYGRCLGFQFTP
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/db_xref="MGD:MGI:96790"
/trans1ation="MEPAVESAPVGAQASKQGKEGSKNRSRRRWRKGKIKASAFSHSM
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/clone_lib="RIKEN full-length enriched mouse cDNA library"
/dev_stage="adult"
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/db_xref="MGD:MGI:1912758"
/db_xref="taxon:10090"
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/strain="C57BL/6J"
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Mammalia; Eutheria; Primates; Craniata; Vertebrata; Eutharyota; Metazoa; Chordata; Craniata; Vertebrata; Euthammalia; Eutheria; Primates; Catarrhini; Hominidae; Howard in the Collection of the
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DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can
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Location/Qualifiers
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/db_xref="taxon:9606"
/clone="NAGE:5744119"
/clone_ib="NHH_MGC_119"
/tissue_type="medulla"
/lab_host="DH10B"
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          (Invitrogen). Research Genethis is a NIH_MGC Library."
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                                                                                                                                                                                                                                                                                                                           Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLN
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information
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NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian
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603077016F1 NIH_MGC_119
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Contact: Robert Strausberg, Ph.D.
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Plate: LLAM11419 row: p column:
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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                                                                                                          /organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5168882"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
/lab_host="DH108"
/note="Organ: brain; Vector: pCMV-SPORT6; Site_1: Not1; Site_2: EcoRV (destroyed); RNA source normal medulla fro anonymous male age 27. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.3 kb, insert size range 0.9-3 kb. Library is normalized and enriched for
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mRNA sequence.
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AGCCTGCTTCAAACCAAAGACCCCTCACCCAGCAGGAGACCCCTGCACAACATGATGCTG
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Clone distribution: MGC clone distribution information
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
plate: LLAM11412 row: m column: 12
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Tissue Procurement: Life Technologies,
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National Institutes of Health, M
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1 (bases 1 to 905)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Ge
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
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B1827706.1 GI:
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                                               /organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5165959"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
/lab_host="DH10B"
/note-"Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI; Site_2: EcoRV (destroyed); RNA source normal medulla fro anonymous male age 27. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.3 kb, insert size range
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        GGATAGAACCTGGAAGATCAATATCTCCCGTGAGGGAAATAACAATGGAGCCAGGTTCTA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Life Technologies, Inc.
CDNA Library Preparation: Life Technologies, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LIN
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information
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1 (bases 1 to 870)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian
Unpublished (1999)
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/clone="IMAGE:5165788"
/clone_lib="NIH_MGC_119"
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/lab_host="DH10B"
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RESULT 8 BI561175 LOCUS SOURCE ORGANISM ACCESSION VERSION FEATURES COMMENT REFERENCE DEFINITION KEYWORDS TITLE AUTHORS JOURNAL DNA Sequencing by: Incyte Genomics, Inc. Clone distribution: MGC clone distribution information found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Plate: LLAM11749 row: g column: 04
High quality sequence stop: 820.
Location/Qualifiers Tissue Procurement: Miklos Palkovits, M.D., Ph.D. cDNA Library Preparation: Michael J. Brownstein (NHGRI), Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 871)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC) Contact: Robert Strausberg, Ph.D. Email: cgapbs-r@mail.nih.gov Unpublished (1999) mRNA sequence. BI561175 603253712F1 NIH\_MGC\_97 Homo sapiens cDNA clone IMAGE:5296155 5' в1561175 Homo sapiens human EST BIS61175.1 /organism="Homo sapiens" GI:15448489 871 bp mRNA EST 05-SEP-2001 Shiraki be

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                                   CTCCAGCTCAACAAGAAGCTGAATCAACACCTGCGGCCCAGGCTAAACCTGGAGCCAAAA 776
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                       CTCCAGCTCAACAAGAAGCTGAATCAACACCTGCGGCACAGGCTAAACCTGGAGCC-AAA
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Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies,
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              CDNA Library Preparation: Life Technologies, Inc. CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Incyte Genomics, Inc. Clone distribution: MGC clone distribution information can found through the I.M.A.G.E. Consortium/LLNL at: http://inage.llnl.gov plate: LLAM11429 row: p column: 09
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National Institutes of Health, Mammalian Gene Collection (MGC).
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//note="Organ: brain; Vector: pCMV-SPORT6; SILE_1. NOTE."
//note="Organ: brain; Vector: pCMV-SPORT6; SILE_1. NOTE."
Site_2: ECORV (destroyed); RNA source normal medulla from anonymous male age 27. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.3 kb, insert size range 0.9-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber full-length clones and was constructed by C. Gruber full-tength clones and Genetics tracking code 013. Note:
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                               Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LLAM11405 row: g column: 01
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                                                                                                                                                                                                                           mRNA sequence.
BI520246
BI520246.1 GI:
EST.
                                                                                                                                                 Bukaryota; Metazoa; Chordata; Craniata; V
Mammalla; Eutherla; Primates; Catarrhini;
1 (bases 1 to 788)
NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian
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603071104F1 NIH_MGC_119
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                             quality sequence stop:
/organism="Homo
                    Location/Qualifiers
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sapiens'
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Catarrhini; Hominidae
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Query Match
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Matches 777
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TGCCATCCGGCCATTCCTGCAGACCATCTCCATTGGGCTGGTGTCCTTCGGGGAGCACTA 1764
                                                                                                                               GTATGTCACGCTGCATAAGGGATGCTTCTATGGCCGCTGCCTGGGCTTCCAGTTCACGCC 1704
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/clone="IMAGE:5163288"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
/lab_host="DH10B"
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Pred. No. 2.3e-160;
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                                                                                                                               TCAAGGCTCATCCACACATGGACCTGCGCACAATGACACAGTCGCTGGTGACTCTGGCG
GGCGTTTTTGCCGGTGTACGGGAGCAGGCGCTGGGGCCTGGAGCCCGGCCCTGGGCCCCTG 1342
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NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian Gene Collection (MGC)
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae,
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Plate: LLAM11405 row: j column: 01
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/clone="IMAGE:5163360"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
/lab_host="DH10B"
                                                                                                                                                                                                                                                                                                                                                                                                                                inote-"Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI; Site_1: EcoRV (destroyed); RNA source normal medulla from anonymous male age 27. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.3 kb, insert size range 0.9-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 013. Note: this is a NHLMGC Library."
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99.5%;
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Pred. No. 3.4e-154;
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Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         BI832284.1 GI:15943834
EST.
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603080125F1 NIH_MGC_119
                                                                                                                                                                                      http://image.llnl.gov
Plate: LLAM11427 row: b column:
                                                                                                                                                                                                                                                                                                                                               Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
                                                                                                                                                                                                                                                                                                                                                                              NIH-MGC http://mgc.nci.nih.gov/.
National Institutes of Health, Mammalian
                                                                                                                                                                                                                                                                                                                                                                                                                                    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          mRNA sequence
                                                                                                                                                                       High quality sequence stop: 781.
                                                                                                                                                                                                                      found through the I.M.A.G.E. Consortium/LLNL
                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Homo sapiens
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                              /organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:5171623"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
/lab_host="DH10B"
/note="Organ: brain; Vector: pCMV-SPORT6; Site_1: NotI;
Site_2: EcoRV (destroyed); RNA source normal medulla from
                                                                                                                                                     ocation/Qualifiers
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                                                                                                                                                                                                                                                                          -TCCCAGCAGGACCTGGGCTAGGAAAAGAATCTATAACTCAACAGGAGCCAGCATTGAGA
AI337272 767 bp mRNA linear EST 18-MAR-1999 tb96h07.xl NCI_CGAP_Col6 Homo sapiens cDNA clone IMAGE:2062237 3' similar to gb:Ll1706_cds1 HORMONE SENSITIVE LIPASE (HUMAN);, mRNA
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        (Invitrogen). Research Genetics tracking code 013. Note: this is a NIH_MGC Library."

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7.9e-151;
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GACAGGACAGTGAGGAGCTCAGCAGCCTGATAAAGTCCAACGGCCAACGGAGCCTGGAGC
                                  GACAGGACAGTGAGGAGCTCAGCAGCCTGATAAAGTCCAACGGCCAACGGAGCCTGGAGC
                                                                       TGGCCCACACAGGCCCTGGGCCCGTCCTCGTCAGGCTCATCTCCTATGACCTGCGTGAAG
                                                                                                                                               AAGCCTTTGAGATGCCACTGACTGCCGACCCCACGCTCACGGTCACCATCTCACCCCCAC
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             CDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome
Clone distribution: NCI-CGAP clone distribution
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 3032 Std Error: 0.00
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Email: cgapbs-r@mail.nih.gov
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AI337272.1 GI:4074199
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National Cancer Institute, Cancer Genome Anatomy
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/lab_host="DH10B"
/note="Organ: colon; Vector: pT7
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/clone_lib="NCI_CGAP_Co16"
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/db_xref="taxon:9606"
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Pred. No. 1.3e-147;
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5 others
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JOURNAL
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Homo sapiens, clone IMAG:
BC029301
BC029301.1 GI:20381052
HTC.
                                                                                                                                                                                                                                                                                                                                                                                                                                                     Submitted (01-MAY-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
                                                                                                Clone distribution: MGC clone distribution information can be fo through the I.M.A.G.E. Consortlum/LINL at: http://image.llnl.gov Series: IRAK Plate: 51 Row: c Column: 24
This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF
                                                                                                                                                                                                                                                                                                                           Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium
DNA Sequencing by: Baylor College of Medicine Human
                                                                                                                                                                                                                        Web site: http://www.hgsc.bcm.tmc.edu/cdna/contact: amg@bcm.tmc.edu
Gontact: amg@bcm.tmc.edu
Gunaratne, P.H., Garcia, A.M., Lu, X., Huly
Yoon, V.S., Kowis, C.R., Lawrence, S., Mart
                                                                                                                                                                                                                                                                                              Sequencing Center
Center code: BCM-HGSC
                                                                                                                                                                                                                                                                                                                                                                                                                   NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
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Mammalia; Eutheria;
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                                                                    clone has the following
/organism="Homo sapiens"
/db_xref="taxon:9606"
                                  Location/Qualifiers
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Primates;
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                                                                    polyA-tail
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ACCESSION VERSION

SOURCE

ORGANISM

Homo sapiens Eukaryota; Metazoa; Mammalia; Eutheria;

Chordata; Primates;

Craniata; Vertebrata; Catarrhini; Hominidae;

Euteleostomi Homo.

KEYWORDS

mRNA sequence. BI826036 BI826036.1 GJ EST. BI826036 LOCUS

DEFINITION

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                                    CTCTCTTCACCAGCGGCCGC 1367
                                                                                        TGGTGTCCTTCGGGGAGCACTACAAACGCAACGAGACAGGCCTCAGTGTGGCCGCCAGCT
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                                                             CTCTCTTCACCAGCGGCCGC 1822
                                                                                                                  TGGTGTCCTTCGGGGAGCACTACAAACGCAACGAGACAGGCCTCAGTGTGGCCGCCAGCT
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/tissue_type="Brain, adult me/clone_lib="NuH_MGC_119"
/lab_host="DH10B"
/note="Vector: pCMV-SPORT6"
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99.0%;
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Best Local :
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            537 TTCTTGCCCCACAGAAGCCCCGCACCACCAGCAATCACCTTACATCCAAAGGGTGCTGCTCA 596
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                                              ATGATGCTGAATCCCAGAAGGAACCTAGAGCCCAACAAAAATCTGCTTCACAAGAGGAAT 480
                                                                   ATGATGCTGAATCCCAGAAGGAACCTAGAGCCCAACAAAAATCTGCTTCACAAGAGGAAT 536
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CTGGGCCAGAAAAGACACCCATAGCCCAGCCAGAATCGAAGACTCTGCAGGGATCCAATA
                                                                                                                                                                                                                                                                                                                                                                                                                                         Email: cgapbs-r@mail.nih.gov
Tlssue Procurement: Life Technologies, Inc.
cDNA Library Preparation: Life Technologies, Inc.
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         http://image.llnl.gov
Plate: LLAM11418 row: a column:
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Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
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NIH-MGC http://mgc.nci.nih.gov/.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    /note="Organ: brain; Vector: pCMV-SPORT6; Site_1: Not1; Site_2: EcoRV (destroyed); RNA source normal medulla from anonymous male age 27. Library is oligo-dry primed and directionally cloned (EcoRV Site is destroyed upon cloning). Average insert size 1.3 kb, insert size range 0.9-3 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 013. Note: this is a NIH_MGC Library."
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             /clone="IMAGE:5168147"
/clone_lib="NIH_MGC_119"
/tissue_type="medulla"
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/db_xref="taxon:9606"
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                                                                                        CTCAACAGGAAGCTGCCTCCCAGCAGGGACCTGGGCTAGGAAAAGAATCTATAACTCAAC 600
                                                                                                     CTCAACAGGAAGCTGCCTCCCAGCAGGAGCACGTGGGCTAGGAAAAGAATCTATAACTCAAC 656
                                                                                                                                       TTCTTGCCCCACAGAAGCCCGCACCACAGCAATCACCTTACATCCAAAGGGTGCTGCTCA 540
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Search completed: January 3, 2003, 07:06:07 Job time: 3339 secs